

WEST Search History

DATE: Thursday, May 06, 2004

Hide?	Set Name	Query	Hit Count
<i>DB=USPT,EPAB,DWPI; PLUR=YES; OP=ADJ</i>			
<input type="checkbox"/>	L1	molecular break light	1
<input type="checkbox"/>	L2	probe same (hairpin or stem loop)	616
<input type="checkbox"/>	L3	probe same (hairpin oligonucleotide or (stem loop near oligonucleotide))	25
<input type="checkbox"/>	L4	L2 and fluorophore	263
<input type="checkbox"/>	L5	L4 and quencher	133
<input type="checkbox"/>	L6	L5 and enediyne	1
<input type="checkbox"/>	L7	L5 and cleavage agent	11
<input type="checkbox"/>	L8	L5 and bleomycin	0
<input type="checkbox"/>	L9	enediyne	156
<input type="checkbox"/>	L10	L9 and probe	55
<input type="checkbox"/>	L11	L10 and fluorophore	11
<input type="checkbox"/>	L12	L11 and cleav\$	11
<input type="checkbox"/>	L13	L7 and cleav\$	11
<input type="checkbox"/>	L14	molecular beacon probes	156
<input type="checkbox"/>	L15	L14 and (bleomycin or enediyne)	2
<input type="checkbox"/>	L16	L14 and L9	1
<input type="checkbox"/>	L17	L14 and (fluorophore and quencher)	69
<input type="checkbox"/>	L18	L17 and (hairpin oligonucleotide or stem loop oligonucleotide)	7
<input type="checkbox"/>	L19	L18 and cleav\$	4
<i>DB=PGPB,USPT,USOC,EPAB,DWPI; PLUR=YES; OP=ADJ</i>			
<input type="checkbox"/>	L20	L14 and cleavage agent	5
<input type="checkbox"/>	L21	L14 and (exonuclease or restriction enzyme or restriction endonuclease or nuclease)	113
<input type="checkbox"/>	L22	L21 and (stem loop or hairpin)	69
<input type="checkbox"/>	L23	L22 and ((fluorophore or acceptor) and quencher)	55
<input type="checkbox"/>	L24	L23 and solid support	30
<input type="checkbox"/>	L25	calicheamicin	396
<input type="checkbox"/>	L26	L25 and cleav\$	331
<input type="checkbox"/>	L27	L26 and L21	0
<input type="checkbox"/>	L28	L26 and (fluorophore and quencher)	14

<input type="checkbox"/>	L29	CalC and l26	10
<input type="checkbox"/>	L30	L29 and l28	1
<input type="checkbox"/>	L31	nucleotd=ide protective agent	0
<input type="checkbox"/>	L32	nucleotide protective agent	2
<input type="checkbox"/>	L33	(protective agent near (nucleic acid or nucleotide))	2
<input type="checkbox"/>	L34	CalC or calicheamicin-resistance	18665
<input type="checkbox"/>	L35	WO 0037608	0
<input type="checkbox"/>	L36	WO0037608	0
<input type="checkbox"/>	L37	WO 0037608	0
<input type="checkbox"/>	L38	WO000037608	0

END OF SEARCH HISTORY

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-24.95	-24.95

FILE 'STNGUIDE' ENTERED AT 14:54:33 ON 06 MAY 2004
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FILE CONTAINS CURRENT INFORMATION.
 LAST RELOADED: Apr 30, 2004 (20040430/UP).

=> d his

(FILE 'HOME' ENTERED AT 14:29:49 ON 06 MAY 2004)

FILE 'MEDLINE, BIOTECHDS, EMBASE, BIOSIS, SCISEARCH, CANCERLIT, CAPLUS'
 ENTERED AT 14:30:22 ON 06 MAY 2004

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L1      2 S (MOLECULAR BREAK LIGHT) AND PROB##
L2      19 S (HAIRPIN OR STEM LOOP) AND MOLECULAR BEACON PROB##
L3      15 DUP REM L2 (4 DUPLICATES REMOVED)
L4      1 S (L3 AND (EXONUCLEASE OR NUCLEASE OR ENDONUCLEASE OR RESTRICT
L5      1 S L3 AND CLEAV?
L6      2642 S (HAIRPIN OR STEM LOOP) AND PROB##
L7      74 S L6 AND (FLUOROPHORE AND QUENCHER)
L8      48 DUP REM L7 (26 DUPLICATES REMOVED)

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FILE 'STNGUIDE' ENTERED AT 14:54:33 ON 06 MAY 2004

=>

Designing a novel molecular beacon for surface-immobilized
DNA hybridization studies;
a DNA **probe** with a **stem loop**
structure that emits fluorescence upon binding to a target
DNA, for use as a DNA biosensor

AUTHOR: Fang X; Liu X; Schuster S; *Tan W

CORPORATE SOURCE: Univ.Florida

LOCATION: Department of Chemistry and UF Brain Institute, University of
Florida, Gainesville, FL 32601, USA.

SOURCE: J.Am.Chem.Soc.; (1999) 121, 12, 2921-22

CODEN: JACSAT

ISSN: 0002-7863

DOCUMENT TYPE: Journal

LANGUAGE: English

AN 1999-05924 BIOTECHDS

AB A biotinylated ssDNA molecular beacon was designed for use in DNA hybridization analysis. DNA hybridization and molecular interaction studies are important techniques for genetic disease diagnosis, particularly where clinical symptoms are linked to nucleic acid mutations. Molecular beacons (MB) are a new class of DNA **probe**, and are used to identify the mutations responsible for genetic disease. MB are ss DNA **probes** containing a **stem loop** structure. The loop structure is capable of reporting the presence of a specific complementary DNA sequence, and the five bases at each end of the MB are complementary, forming a stem. A **quencher** is immobilized to one of the ends, and a **fluorophore** to the other, so that the fluorescence is quenched until the loop comes into contact with a target DNA sequence. The hybrid between loop and target DNA is longer and more stable than the stem structure, and thus the stem is broken, and the **fluorophore** brought out of contact with the **quencher**, resulting in fluorescent emissions. This is of significant value in studying genetics, particularly genetic disease and molecular interaction. It can also be used to produce DNA biosensors.
(14 ref)